

:. ADA 039537

UNCLASSIFIED SECURITY CLASSIFICATION OF THIS PAGE (When Date Entered) READ INSTRUCTIONS
BEFORE COMPLETING FORM REPORT DOCUMENTATION PAGE RECIPIENT'S CATALOG NUMBER 2. GOVT ACCESSION NO. HDL-TR-1806 TYPE OF REPORT & PERIOD COVERED Charge Yield and Dose Effects in MOS Technical Report Capacitors at 80 K . B. PERFORMING ORG. REPORT NUMBER 8. CONTRACT OR GRANT NUMBER(*) Harold E. Boesch, Jr. James M./McGarrity PERFORMING ORGANIZATION NAME AND ADDRESS 10. PROGRAM ELEMENT, PROJECT, TASK AREA & WORK UNIT NUMBERS Harry Diamond Laboratories 2800 Powder Mill Road Program Ele: 6.27.04.H Adelphi, MD 20783
11. CONTROLLING OFFICE NAME AND ADDRESS 77 Director Defense Nuclear Agency Washington, DC 20301
MONITORING AGENCY NAME & ADDRESS(II dilierent from Controlling Office) 15. SECURITY CLASS. Unclassified 15a. DECLASSIFICATION/DOWNGRADING 16. DISTRIBUTION STATEMENT (of this Report) Approved for public release; distribution unlimited. 17. DISTRIBUTION STATEMENT (of the abstract entered in Block 20, If different from Report) 63 050 SUPPLEMENTARY NOTES
HDL Project: 236T28 DRCMS Code: 6970002211556 This work was sponsored by the U.S. Army Materiel Development and Readiness Command and the Defense Nuclear Agency under Z99QAXT007, Work Unit 64, Unit Title "TRANSEFF." 19. KEY WORDS (Continue on reverse side if necessary and identify by block number) MOS capacitors Hole transport Low temperature Radiation damage Charge yield D. ABSTRACT (Continue on reverse side if necessary and identity by black number) The response of metal-oxide semiconductor capacitors to shortpulse high-energy electron irradiation was measured at 80 K. It was confirmed that radiation-generated holes are almost totally retained in the SiO₂ layer regardless of oxide processing; this retention was exploited to determine carrier yield as a function of applied electric field. Evidence was found that the holes

DD 1 JAN 73 1473 EDITION OF 1 NOV 65 IS OBSOLETE

UNCLASSIFIED

1 SECURITY CLASSIFICATION OF THIS PAGE (When Date Entered)

over

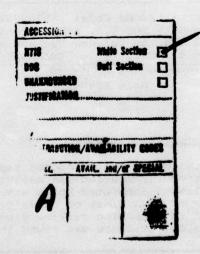
bpg

SECURITY CLASSIFICATION OF THIS PAGE(When Date Entered)

1,000,000

undergo an initial displacement under an applied field immediately following carrier generation (9.5 nm at $10^5 \, \text{V/cm}$). Samples were subjected to pulsed irradiation at 80 K to accumulated doses above $10^5 \, \text{rads}(\text{SiO}_2^0)$ and mechanisms which limit hole buildup above $5 \times 10^6 \, \text{rads}(\text{SiO}_2^0)$ were explored. Electron-hole recombination in a low field region of the SiO was identified as an important process and was modeled. Other mechanisms discussed include electron injection, field—and temperature—activated hole transport, applied field collapse, and dielectric breakdown.

50,000



CONTENTS

di

		Page
1.	INTRODUCTION	5
2.	SAMPLES AND EXPERIMENTAL TECHNIQUES	6
3.	CHARGE-YIELD MEASUREMENT	7
	3.1 Field Dependence of Hole Yield	7
	3.2 Initial Hole Displacement	10
4.	DOSE DEPENDENCE OF CHARGE BUILDUP	13
	4.1 Measurements and Results	13
	4.2 Discussion	15
	4.2.1 Linear Region	15 16
	4.2.2.1 Recombination in Low-Field Region	16
	4.2.2.2 Electron Injection at High Fields	21
	4.2.2.3 Hole Transport	22
	4.2.2.4 Collapse of Applied Field	23
	4.2.2.5 Dielectric Breakdown	24
	4.2.2.6 Applicability to Higher Temperatures	24
5.	CONCLUSIONS	25
	ACKNOWLEDGEMENTS	25
		25
	LITERATURE ĆITED	25
	DISTRIBUTION	27
	FIGURES	
	1 Flatband-voltage shift and corresponding hole yield in MOS capacitors irradiated at 80 K	9
	2 Normalized flatband-voltage shift and equivalent	
	flatband shift for E = 1 MV/cm and L = 100 nm for various MOS capacitors at 80 K	14
	3 Flatband-voltage shift, ΔV_{FR} , 1 ms after irradiation of	
	HAC p-type and n-type MOS capacitors at 80 K	14

CONTENTS (Cont'd)

		Page
4	Flatband-voltage shift, $\Delta V_{\rm FB},$ as function of time after first LINAC pulse delivered to HAC dry-n MOS capacitors .	15
5	Model for limitation of charge buildup in an oxide layer	17
6	Calculated field and hole density profiles in the oxide as a function of dose at 80 K	20
7	Comparison of model calculations and experimental results on the flatband-voltage shift	20
	TABLES	
I	MOS Capacitor SiO ₂ Characteristics	6
II	Initial Hole Displacement in MOS Samples at $\sim 80~K$	11

1. INTRODUCTION

Metal-oxide semiconductor (MOS) devices, such as charge-coupled devices (CCD's) and MOSFET's (MOS field-effect transistors), are being investigated for use in imaging, signal processing, and detector preamplifier applications in which the detector and its immediately associated electronics are operated at low temperatures (<80 K). Recent experiments on MOS capacitors and devices at ~80 K have established that, when hole-electron pairs are generated in the SiO2 insulating layer by ionizing radiation, the electrons are rapidly swept out of the oxide while the holes remain essentially immobile ("frozen in") at or near their point of creation. 1, 2, 3 As a result, most of the holes generated by the radiation remain in place in the oxide layer for significant times and cause relatively large flatband- or threshold-voltage shifts per unit dose in MOS structures. (A flatband voltage shift of ~-1 V is expected for a 10-krad(SiO₂) dose in a 100-nm oxide.) These shifts threaten device operation at doses well below 100 krads. In contrast to the highly process-dependent "permanent" trapping of a small fraction (typically 1 to 10 percent) of the radiation-generated holes observed at room temperature in "hard" MOS oxides, the retention of the holes in SiO2 at ~80 K is largely process independent2 and results from a strongly temperature-dependent hole-transport mechanism. 1, 3, 4 Since the yield of electron-hole pairs generated in an MOS oxide per unit radiation dose is believed⁵ to be an intrinsic property of amorphous Sio2, this yield also should be process independent. Therefore, if MOSFET's, MOS integrated circuits (IC's), or charge-coupled devices with pure SiO2 gate-insulator layers are operated and irradiated at low temperatures, they will suffer relatively large oxide-charge buildups whether or not they have been hardened to charge trapping at room temperature.

¹H. E. Boesch, Jr., F. B. McLean, J. M. McGarrity, and G. A. Ausman, Jr., IEEE Trans. Nucl. Sci. NS-22, 2163 (1975).

²H. H. Sander and B. L. Gregory, IEEE Trans. Nucl. Sci. NS-22, 2157 (1975).

³R. C. Hughes, E. P. Eer Nisse, and H. J. Stein, IEEE Trans. Nucl. Sci. NS-22, 2227 (1975).

⁴F. B. McLean, H. E. Boesch, Jr., and J. M. McGarrity, Hole Transport and Recovery Characteristics of MOS Gate Insulators, IEEE Nuclear and Space Radiation Effects Conference (July 1976).

⁵G. A. Ausman, Jr., and F. B. McLean, Appl. Phys. Lett. <u>26</u>, 173 (1975).

The present report discusses two related investigations of radiation-induced charge buildup in SiO_2 at ~ 80 K. In the first study, the retention of holes at low temperature was exploited to provide a direct measure of hole-electron pair yield per unit radiation dose at low temperature as a function of electric field for high-energy ionizing radiation. In the second study, the response of MOS capacitors to short-pulse high-dose irradiation at ~ 80 K was investigated, and mechanisms which impose limits on the oxide-charge buildup were examined.

2. SAMPLES AND EXPERIMENTAL TECHNIQUES

Dry-grown oxide samples were provided by Hughes Aircraft Corporation (HAC), Northrop Research and Technology Center (NRTC), and a proprietary source supported by the Defense Nuclear Agency (DNA). HAC also supplied a wet-oxide sample representative of the process under development in their hardened complementary-MOS/silicon-on-sapphire program. The dry oxides were thermally grown at 1000°C and the wet oxide was pyrogenically grown at 950°C, usually on n-type Si (see table I). Aluminum gate electrodes were deposited, and the resulting MOS capacitors were bonded to headers without caps.

TABLE I. MOS CAPACITOR SIO, CHARACTERISTICS

Identifier	Gate deposition	Post anneal	Oxide thickness (nm)
DNA No. 1	filament	A, 15 m, 800°C	72.5
DNA No. 2	filament	None	65.5
HAC dry n	C crucible	None	87.5
HAC dry p	C crucible	None	87.5
HAC wet n	C crucible	N ₂ , 20 m, 925°C	96.5
NRTC 850	e-beam	None	85
NRTC 671	e-beam	None	67
NRTC 1538	e-beam	None	154

The experiments were performed with the electron linear accelerator (LINAC) at the Armed Forces Radiobiology Research Institute (AFRRI). The LINAC produced a nominal 12- to 13-MeV 1-A electron beam with a pulse width of 4 us. Multiple pulses could be delivered to the sample at a rate of 60/s. Sample dose per pulse was controlled by varying the LINAC-to-sample distance. A thin-foil Cu calorimeter was used for pulse-to-pulse electron-beam dosimetry. Teflon:CaF2 thin-disc thermoluminescent dosimeters were employed for absolute dose measurements. For the charge-yield experiment, the samples were mounted in a liquid-nitrogen-cooled sample holder under vacuum. 6 For the high-dose experiments, the samples were submerged directly in a liquid-nitrogen bath to insure maximum thermal transfer. For both experiments, data were taken only on the first pulse delivered to a sample. A fast high-frequency capacitance-voltage (C-V) measuring apparatus6,7 recorded preirradiation C-V characteristics and monitored characteristics as a function of time either after a single radiation pulse, or, sometimes, at 1 and 12 ms after each individual radiation pulse in a multiple-pulse burst. The MOS capacitance of the sample was monitored by a phase-sensitive detector system operating at 5 MHz. The C-V characteristics were recorded on oscilloscopes by monitoring the capacitance as a 0.1 ms voltage ramp was applied to the capacitor. The fast C-V system was intercalibrated with a Boonton 71A capacitance meter and found capable of accurately reproducing the deep-depletion C-V trace. From the C-V data, the radiation-induced flatband-voltage shift, ΔV_{FR} , was extracted.

3. CHARGE-YIELD MEASUREMENT

3.1 Field Dependence of Hole Yield

The field (bias) dependence of the yield of charge carriers in \sin_2 was measured by Curtis, Srour and Chiu⁸ at room temperature by a charge-collection technique and low-energy (~4-keV) electron irradiation. Ausman and McLean⁵ demonstrated that these results were consistent with a hole-electron pair-creation energy W ~18 eV/pair,

⁵G. A. Ausman, Jr., and F. B. McLean, Appl. Phys. Lett. <u>26</u>, 173 (1975).

⁶H. E. Boesch, Jr., Development of Apparatus for Performing Rapid Capacitance-Voltage Measurements on MIS Structures, Harry Diamond Laboratories TM-76-33 (December 1976).

⁷F. B. McLean, H. E. Boesch, Jr., P. S. Winokur, J. M. McGarrity, and R. B. Oswald, Jr., IEEE Trans. Nucl. Sci. NS-21, 47 (1974).

⁸O. L. Curtis, Jr., J. R. Srour, and K. Y. Chiu, J. Appl. Phys. <u>45</u>, 4506 (1974).

which in turn agrees with results of a model for the ionization process based on plasmon creation. Following creation, the actual yield of free carriers is determined by field-aided escape (i.e., aided by applied bias) of carriers from bimolecular recombination in the high ionization-density regions along the tracks of the incident kilovolt electrons. Sonowden et al measured collected charge in SiO_2 capacitors at room temperature using high-energy ionizing radiation (30-MeV LINAC electrons). Their results are again consistent with $W_O = 18$ eV/pair. In contrast to kiloelectron volt electron irradiation, which produces high ionization-density regions along the relatively short tracks of the incident particles, the very high energy LINAC electron beam produces widely dispersed point ionization in the SiO_2 similar to that which would be produced by energetic photon irradiation.

In this experiment, hole yield in the SiO_2 induced by 13-MeV LINAC radiation pulses was measured at low temperature by the fast C-V measurement technique. Bias voltages were applied to produce fields from -0.6×10^6 to 4.7×10^6 V/cm. To insure that the radiation-generated oxide charge did not significantly perturb the externally applied oxide field, the dose per radiation pulse was maintained below 3 \times 10 rads(SiO₂).

The early (~l ms after pulse) flatband voltage shift ΔV_{FB} is plotted in figure 1 as a function of oxide internal field for the DNA No. 1 n-type samples. (Similar results were obtained for NRTC 1538A MOS capacitors.) To correct for pulse-to-pulse dose variations, the ΔV_{FB} values have been normalized by the calorimeter readings to a nominal dose of 2 × 10 3 rads(SiO2). The flatband shift saturates beyond 10^6 V/cm and is essentially symmetrical about the zero V/cm axis.

For a uniform radiation-produced oxide charge density, $\rho\,,$ consisting of the holes only,

$$\Delta V_{FB} = -\frac{\rho L^2}{2\varepsilon} \tag{1}$$

⁵G. A. Ausman, Jr., and F. B. McLean, Appl. Phys. Lett. <u>26</u>, 173 (1975).

⁹R. E. Leadon, D. P. Snowden, and J. M. Wilkenfeld, Radiation Effects in Semiconductor and Insulator Materials, IRT Corporation, Harry Diamond Laboratories CR-76-152-1 (April 1976).

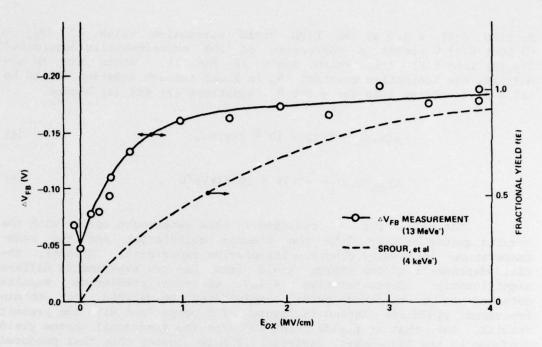


Figure 1. Flatband-voltage shift and corresponding hole yield in MOS capacitors irradiated at 80 K.

where L is the oxide thickness in centimeters and ϵ is the dielectric constant in F/cm. The charge density may in turn be expressed as a function of oxide dose D, in rads(SiO₂), using appropriate material parameters and the electric-field-dependent fraction f(E) of the radiation-generated holes which escape initial recombination,

$$\rho(D,E) = 2.2 \times 10^{-5} \text{ W}_{0}^{-1} \text{ f(E)D}$$
 (2)

Then

$$\Delta V_{FB}(D,E) = (-3.23 \times 10^7) \text{ f(E)} W_0^{-1} L^2 D$$
 (3)

The fractional hole yield as a function of field, f(E), may be obtained from the ΔV_{FB} data in figure 1 by assuming that the asymptotic or saturation value of the flatband shift above $\sim 3 \times 10^6$ V/cm corresponds to all the holes escaping initial recombination; i.e., f(E) = 1.0.

Setting f(E) = 1.0 at the high field saturation value of $\Delta V_{\rm FB}$ = -0.18 ± 0.03 V yields a conversion of the experimentally determined $\Delta V_{\rm FB}(E)$ into f(E) (right-hand scale in fig. 1). Since f(E) is now defined, the ionization constant $W_{\rm O}$ is found through equation (3) to be (18 ± 3)eV. Using this value for $W_{\rm O}$, equations (2) and (3) become

$$\rho(D,E) = 1.22 \times 10^{-6} f(E)D$$
, (4)

$$\Delta V_{FB}(D,E) = -1.79 \times 10^6 f(E) L^2 D$$
 (5)

The value for W obtained in this experiment agrees with the results quoted earlier from the plasmon calculation and the room-temperature low-energy electron-irradiation experiment. However, the field dependence of the charge yield from the two experiments differs significantly. Representative 4-keV electron-irradiation results obtained on a hardened oxide sample 10 and normalized to obtain fractional yield are plotted in figure 1 for comparison with the present results. Note that at fields below 106 V/cm the fractional charge yield produced by the high-energy radiation is much greater than that produced by the low-energy electrons. The present results are adequately explained by a geminate recombination process in which field-dependent rapid recombination takes place between the members of the widely dispersed electron-hole pairs produced by the high energy radiation.*

3.2 Initial Hole Displacement

To this point, it has been assumed that all the holes remain in the oxide after irradiation. However, evidence has been found in this work that a fraction of the holes escapes the oxide immediately following generation, even at $\sim\!80~\rm K$. As indicated in a previous paper, the flatband shifts obtained in identical MOS samples at identical doses under negative and positive biases are almost, but not quite, the same. Results obtained on a number of samples from a pair of such measurements are recorded in table II. In each case, the negative bias shift is 10 to 20 percent less than the positive shift. Initially,

^{*}G. A. Ausman, Jr., private communication.

¹H. E. Boesch, Jr., F. B. McLean, J. M. McGarrity, and G. A. Ausman, Jr., IEEE Trans. Nucl. Sci. NS-22, 2163 (1975).

¹⁰J. R. Srour, O. L. Curtis, Jr., and K. Y. Chiu, IEEE Trans. Nucl. Sci. NS-21, No. 6, 73 (1974).

TABLE II. INITIAL HOLE DISPLACEMENT IN MOS SAMPLES AT ~80 K

Sample	△VFB+ (V)	∆VFB- (V)	E (MV/cm)	Distance (nm)
DNA No. 1	-3.5	-2.7	1.38	9.4 ± 1.4
DNA No. 2	-3.3	-2.5	1.53	9.0 ± 1.3
HAC dry n	-2.7	-2.1	1.14	10.9 ± 2.1
NRTC 850	-6.2	-5.0	1.18	9.1 ± 0.8
NRTC 1538	-2.7	-2.3	0.98	12.3 ± 3.3

it was assumed that this variation resulted from a decrease in the oxide field under negative bias because of a potential drop across the depletion region in the n-type silicon under inversion or deep-depletion conditions. However, this potential drop virtually disappears during a radiation pulse since the depletion region collapses very quickly as carriers are generated in the silicon and transported through the depletion region. Also, the $\Delta V_{\rm FB}$ relative decrease at negative biases was exhibited by a pair of samples—HAC dry n and HAC dry p—with identical oxide layers which were irradiated at positive and negative biases respectively, i.e., under accumulation conditions.

The bias effect is adequately explained by postulating that the holes are first transported through extended states in the valence band imediately after the holes are thermalized and before they are captured at a defect site and hopping transport begins. If at a given field the holes are transported an average distance d in the valence band in an oxide of thickness L, then under positive bias and uniform field conditions in the oxide, the hole distribution will be uniformly displaced toward the Si. This displacement leaves the zone between x = 0 (where x is a distance from the metal electrode) and x = d depleted of holes. Similarly, under negative bias a zone between x = L - d and x = L (the Si interface) will be depleted. The corresponding $\Delta V_{\rm FB}$ values are

$$\Delta V_{FB+} = -\rho/\epsilon \int_{d}^{L} x dx = -\rho/2\epsilon (L^2 - d^2) , \qquad (6)$$

⁹R. E. Leadon, D. P. Snowden, and J. M. Wilkenfeld, Radiation Effects in Semiconductor and Insulator Materials, IRT Corporation, Harry Diamond Laboratories CR-76-152-1 (April 1976).

$$\Delta V_{\text{FB-}} = -\rho/\epsilon \int_{0}^{\text{L-d}} x dx = -\rho/2\epsilon (\text{L - d})^{2}.$$
 (7)

Then

$$f_{d} = \begin{vmatrix} \Delta V_{FB+} - \Delta V_{FB-} \\ \Delta V_{FB+} \end{vmatrix} = \frac{2d (L - d)}{L^{2} - d^{2}}.$$
 (8)

Solving for d,

$$d = \frac{Lf_d}{2 - f_d} \tag{9}$$

Values for the initial hole displacement d obtained for several of the MOS samples are presented in table II together with error limits calculated from the estimated probable error in the Δv_{FB} values. The agreement within the error limits implies strongly that the charge-displacement analysis is essentially correct and that d ~ 9.5 nm is appropriate for all the materials examined. If d is assumed to be a schubweg, i.e., d = $\mu_H \tau_H E$ where μ_H is the intrinsic hole mobility, τ_H is the hole dwell time in the valence band, and E is the electric field, then the product $\mu_H \tau_H \approx 7 \times 10^{-3}$ V/cm under these conditions. Unfortunately, the measurements were not performed over a sufficient range of electric field to determine whether d is a schubweg or perhaps a field-independent mean free path.

The error, η , introduced by charge displacement in the measurement of total hole yield under positive bias, is easily found from an extension of the analysis above to be

$$\eta = (\rho L^2/\epsilon - \Delta V_{FB+})/(\rho L^2/\epsilon)$$

$$= d^2/L^2 . \qquad (10)$$

For a 72.7-nm oxide and d = 9.5 nm, η = 0.017. Therefore, initial charge displacement had a negligible effect on the charge-yield results (fig. 1).

4. DOSE DEPENDENCE OF CHARGE BUILDUP

4.1 Measurements and Results

The response of MOS capacitors to pulsed irradiation at 80 K was measured as a function of dose, applied bias, and time after irradiation.

In the first experiment, a variety of MOS samples (table I) were irradiated at various biases to doses in the 10^3 to 10^6 rad(SiO₂) range and ΔV_{FB} was recorded at 1 ms after the radiation pulse.

Figure 2 shows the flatband shifts as a function of dose, with ΔV_{FB+} normalized by the fractional charge yield, f(E), from figure 1, and the geometric oxide-thickness dependence. The data points indicate a linear variation of normalized ΔV_{FB} with dose up to ${\sim}10^5$ rads(SiO $_2$).

In the second experiment, HAC MOS samples were subjected to high-dose multiple-pulse irradiation (up to 5 pulses at a rate of 60/s; typically 1.75×10^5 rads(SiO) per pulse). HAC n-type samples were biased at 5, 10, and 20 V during irradiation, while HAC p-type samples were biased at the corresponding negative biases. Flatband shifts were measured at 1 and 12 ms after each radiation pulse. The results are plotted in figure 3. The dashed lines represent the predicted flatband shift based on equation (5) for f(E) = 1.0. Above 10^5 rads(SiO₂) the flatband shift evidently ceases to increase linearly with dose. Under negative bias in particular, ΔV_{FB} saturates strongly at a value near the magnitude of the applied bias voltage. Under positive bias, ΔV_{FB} continues to increase with dose, but at a sublinear rate. These experimental results are in qualitative agreement with the observations of Nielsen and Nichols, 11 who measured charge buildup at 90 K under cobalt-60 irradiation.

In the third experiment, the flatband shift in HAC n-type MOS capacitors was observed as a function of time up to 800 s after multiple-pulse irradiations to total doses from 40 to 875 krads(SiO₂) and at bias voltages from 5 to 20 V. Typical results are plotted in figure 4. The observed early (4 ms) flatband shifts conform well to predictions of the initial shift based on equation (5). Also, the shift observed following a single 40-krad (SiO₂) pulse (curve E) shows little change with time. At higher doses--150 krads(SiO₂) and above--significant decay of the flatband shift occurs, particularly at the higher bias voltages.

 $^{^{11}}R$. L. Nielsen and D. K. Nichols, IEEE Trans. Nucl. Sci. NS-20, No. 6, 319 (1973).

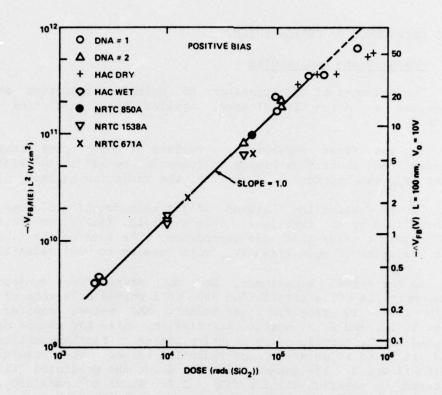


Figure 2. Normalized flatband-voltage shift (left scale) and equivalent flatband shift for $E=1 \, MV/cm$ and $L=100 \, nm$ for various MOS capacitors at 80 K.

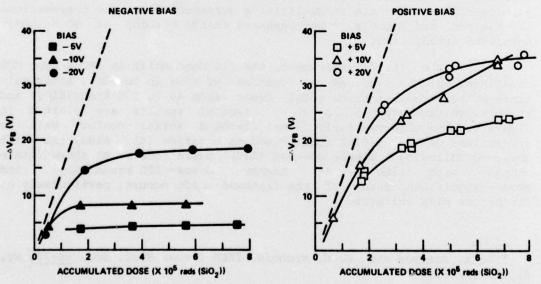


Figure 3. Flatband-voltage shift, $\Delta V_{\rm FB}$, 1 ms after irradiation of HAC p-type (left) and n-type (right) MOS capacitors at 80 K.

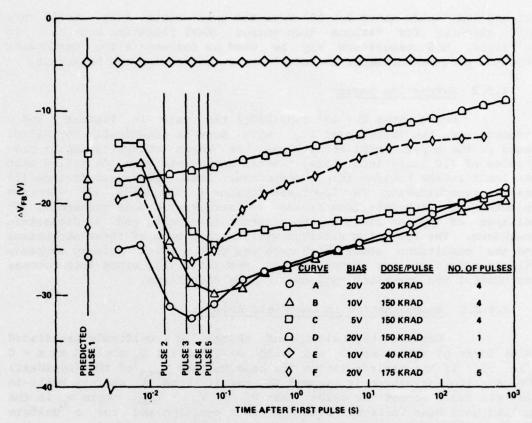


Figure 4. Flatband-voltage shift, ΔV_{FB} , as function of time after first LINAC pulse delivered to HAC dry-n MOS capacitors. Samples received multiple pulses at 60 pulses/s (see legend).

4.2 Discussion

4.2.1 Linear Region

The ΔV_{FB} dose-dependence results presented in figure 2 indicate that, below $\sim\!10^5$ rads(SiO_2), the charge buildup in a variety of SiO_2 samples is properly predicted by equation (5). The 80-K shifts are again shown to be large ($\sim\!-10$ V at 10^5 rads(SiO_2) for 100-nm oxide) and independent of sample source and processing details. This uniform ΔV_{FB} response to pulsed radiation at low temperatures obtained with an early C-V measurement suggests that such capacitors be used as absolute dosimeters. Simply inverting equation (5) yields

$$D = 5.57 \times 10^{-7} \Delta V_{FB+}/f(E)L^{2} . \qquad (11)$$

For applied fields above 2 \times 10 6 V/cm the hole yield f(E) should vary only slightly for various high-energy (MeV) radiation sources. In principle, MOS capacitors may be used as dosimeters for performing absolute dose measurements in actual device geometries and materials.

4.2.2 Saturation Region

Above about 2 \times 10⁵ rads(SiO₂) the data in figures 2 and 3 indicate that the increase of ΔV_{FB} with dose is drastically curtailed. Based on the experimental results and the known charge-transport properties of SiO₂ gate insulators, several mechanisms were identified that can limit charge buildup in MOS structure. These processes include (1) charge recombination in low-field regions in the oxide, (2) electron injection at high-interface fields, (3) accelerated hole transport, (4) collapse of applied field due to electron transport, and (5) dielectric breakdown. The following sections will discuss each of these mechanisms and the conditions under which each may play a role. Primary emphasis will be placed on the recombination mechanism (1), since this process may control MOS structure response in many situations.

4.2.2.1 Recombination in Low-Field Regions

Consider the buildup of charge in a uniformly irradiated oxide layer of thickness L and with a potential V_0 applied at x=0 (fig. 5). If the preirradiation flatband voltage V_{FBO} of the (nonideal) MOS capacitor structure is assumed to result from a uniform built-in electric field across the oxide, then $V_0 = V_G - V_{FBO}$, where V_G is the applied gate bias voltage. From Poisson's equation and for a uniform charge buildup, the electric field in the oxide is given by

$$E(x) = \rho/\varepsilon (x - L/2) + V/L . \qquad (12)$$

For $\rho < |2\epsilon V/L^2|$, corresponding to $|\Delta V_{FB}| < |V_0|$, the electric field in the oxide is monotonic and the radiation-generated electrons are quickly swept out without interacting with the stationary holes (fig. 5a). Note, however, that the electric field is increasing at x=L and decreasing at x=0 because of the contribution of the hole space-charge field. At a critical dose D_{Sat} , ρ approaches $|2\epsilon V_0/L^2|$ (corresponding to the magnitude of the flatband shift approaching the magnitude of the applied potential), and the electric field at x=0 reaches zero and tries to go negative (fig. 5b). In this case, the electrons being swept toward x=0 come to a virtual halt in the zero field region and undergo efficient recombination with the holes in the vicinity. The result is that the hole density near x=0 is rapidly wiped out. As irradiation of the oxide continues (fig. 5c), the zero point for the electric field and, consequently, the zone of rapid recombination move deeper into the

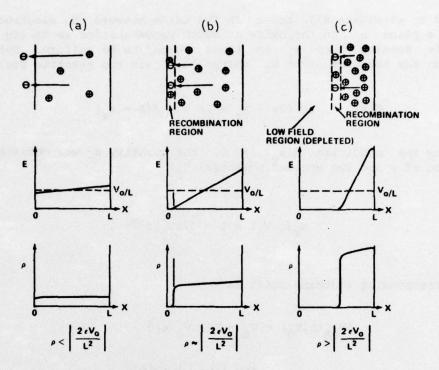


Figure 5. Model for limitation of charge buildup in an oxide layer: (a) electron sweepout; (b) recombination at the interface; (c) recombination in oxide interior.

oxide, "eating away" at the hole distribution. The oxide charge and the total electric field continue to increase at x = L. For irradiation under positive bias, x = 0 in figure 5 corresponds to the electrode interface and x = L is the Si interface. In this case, the recombination process reduces the hole density in a region well removed from the Si interface; consequently, ΔV_{FB+} continues to increase as a function of dose following the onset of recombination, but begins to saturate. For irradiation under negative bias, x = 0 in figure 5 corresponds to the Si interface, and the onset of recombination drastically affects any further increase in ΔV_{FB-} with added dose. In fact, since the net electric field at the Si interface is pinned at zero for $\rho \geq \left|2\epsilon V\right|/L^2$, the flatband shift under negative bias is similarly pinned to ΔV_{FB-} = V_0 . The derivation of the expected dependence on dose of the flatband shift, ΔV_{FB+} under positive bias, was based on this simple model for limiting by recombination and also on certain assumptions. At a given dose above D the recombination process is

assumed to eliminate all holes in the oxide between the electrode (x = 0) and a plane x in the oxide at which recombination is taking place. The hole density from x to L was assumed to be uniform. Poisson's equation for this situation was solved to obtain the electric field:

$$E(x) = \rho/2\varepsilon (2x - x_r - L) + V_0/(L - x_r)$$
 (13)

Imposing the condition $E(\mathbf{x}_r) = 0$, the quantity \mathbf{x}_r was obtained as a function of ρ and the applied potential V_0 :

$$x_r(\rho, V_0) = L - (2\epsilon V_0/\rho)^{\frac{1}{2}}$$
 (14)

The corresponding flatband shift is

$$\Delta V_{FB}(\rho, V_0) = V_0 - L(2\rho V_0/\epsilon)^{\frac{1}{2}}$$
for $|\Delta V_{FB+}| > |V_0|$ (15)

Using equation (4), we may express ΔV_{FB} as a function of oxide dose:

$$\Delta V_{FB+}(D, V_0) = V_0 - 2.68 \times 10^3 L [V_0 f(E) D]^{\frac{1}{2}}$$
for $|\Delta V_{FB+}| > |V_0|$. (16)

This relationship predicts, therefore, that the flatband shift will increase with the square root of the dose for total flatband shifts greater than the applied potential.

The simple recombination model did not include such factors as the field dependence of charge yield in the oxide and the initial displacement of the holes. To aid in a more careful analysis of the $\Delta V_{\mbox{\scriptsize FB}}$ limitation mechanisms, a computer program was written to simulate the charge-buildup process in a basic MOS structure. This

program repetitively solved Poisson's equation in the x dimension as electron-hole pairs were generated in the oxide to obtain the charge and electric field distribution across the oxide layer as a function of time and dose. The generation rate for electron-hole pairs in each x subinterval was governed by the instantaneous electric field in that interval through the experimental yield/field relationship in figure 1. The program allowed for initial displacement of the holes according to the schubweg model, following which the holes were assumed to be immobile. In keeping with the discussion above, the electrons were assumed to be infinitely mobile; that is, they move without limit unless stopped at a point of field reversal. Hole-electron recombination was allowed only at field reversal points.

Representative results from this computer model are shown in figure 6. No parameters were adjusted to obtain these results; all the constants used in the calculation were derived from material properties and the data in figure 1 and table II. Field and hole density profiles in the oxide are shown for doses to 10⁶ rads(SiO₂) for an 87.5-nm oxide under 10-V bias (HAC dry n or p). Note the apparent concentration of the hole buildup and, consequently, the electric field near the x = L interface (Si interface under positive bias) as the dose increases. Other workers have also found evidence for concentration of the holes near the Si interface under positive bias at low temperature. 11,12 It is important to recognize that bunching of the holes near the interface at low temperature does not require hole transport or preferential trapping in that region, but results from the progressive elimination of holes in the balance of the oxide.

Both the simple model and computer model predictions for the flatband shifts in the HAC samples as a function of dose under negative and positive bias are compared in figure 7. Also plotted are the experimental $\Delta V_{\rm FB}$ data from figure 3 (points). Both the simple recombination model and the computer simulation results reproduce the sharp limitation of $\Delta V_{\rm FB}$ to essentially the applied negative bias. The computer calculation shows a stronger limitation of the positive bias shift than predicted by the simple analytic model; this is primarily a result of hole loss by initial transport which becomes significant as the electric field near the Si interface increases and the hole region shrinks. The 5- and 10-V computer-model curves agree with the

 $^{^{11}}R.$ L. Nielsen and D. K. Nichols, IEEE Trans. Nucl. Sci. NS-20, No. 6, 319 (1973).

¹²E. Harari, S. Wang, and B. S. H. Royce, J. Appl. Phys. 46, 1310 (1975).

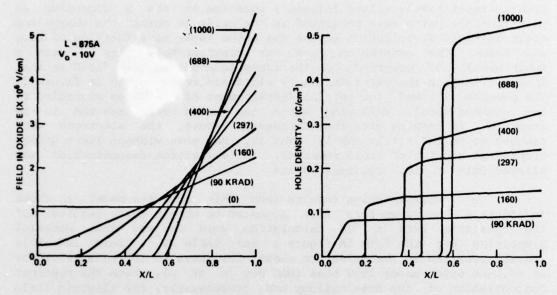


Figure 6. Calculated field and hole density profiles in the oxide as a function of dose at 80 K.

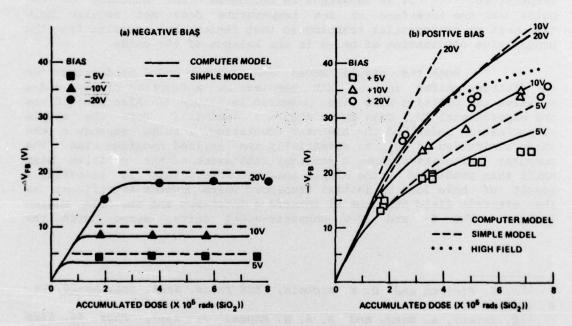


Figure 7. Comparison of model calculations and experimental results on the flatband-voltage shift under (a) positive and (b) negative bias at 80 K.

experimental results, indicating that the recombination mechanism accounts for most of the charge-buildup limitation in these cases. The 20-V bias data points show strong $\Delta V_{\rm FB}$ saturation above 5×10^5 rads(SiO_2) which is not predicted by the computer model; evidently other processes, which will be discussed further, contribute to buildup limitation in this case.

4.2.2.2 Electron Injection at High Fields

The computer simulation of the 20-V buildup predicted that the total electric field at the Si interface should have increased beyond 5 × 10^6 V/cm for doses above 3 × 10^5 rads(SiO₂), while the field in the 10-V case was predicted to remain below that value to above 8 × 10^5 rads(SiO₂). Mitchell¹³ noted that high interface fields would be expected to cause injection of electrons into the oxide from the silicon which would in time result in elimination of some holes in the oxide by recombination. Powell¹⁴ measured significant electron injection via tunneling for interface fields greater than 6 × 10^6 V/cm. Since the tunneling current increased exponentially with the field, this process could result in a strong saturation of the flatband shift above a critical value. A crude attempt at including the electron injection process in the computer model yielded results (dotted line, fig. 7b) in reasonable agreement with the experimental data.

If it is assumed that a mechanism such as tunneling limits the interface field to $E(L) = E_{\max}$, then the maximum flatband shift which can be obtained is

$$\Delta V_{FB+} = V_0 - LE_{max} . (17)$$

The dose at which this flatband shift is first attained may be estimated from the analytic recombination model introduced earlier. Substituting equation (14) into equation (13), setting $E(L) = E_{max}$ and rearranging,

$$\rho\left(E_{\text{max}},V_{0}\right) = \varepsilon E_{\text{max}}^{2}/2V_{0} \qquad (18)$$

or, applying equation (4),

$$D_{\text{sat}} = 1.39 \times 10^{-7} E_{\text{max}}^2 / f(E) V_0$$
 (19)

¹³J. P. Mitchell, IEEE Trans. Elec. Dev. ED-14, No. 11, 764 (1967).
14R. J. Powell, IEEE Trans. Nucl. Sci. NS-22, 2240 (1975).

For an 87.5-nm oxide biased at 20 V and $E_{max}=6.5\times10^6$ V/cm, the saturation shift of -37 V should be attained at 3.7 × 10⁵ rads(SiO₂). For 5-V bias under similar conditions, $\Delta V_{FB+}=-52$ V at D = 1.9 × 10⁶ rads(SiO₂). Thus, field limiting is much more likely to occur at higher applied voltage.

4.2.2.3 Hole Transport

As noted previously, figure 4 shows that significant decay of ΔV_{FB} occurs at 80 K for pulsed irradiation of SiO₂ samples to doses above 150 krads(SiO $_2$). As plotted, ΔV_{FB} for most of the samples decreases linearly as a function of log t (time). This logarithmic flatband shift decay is typical of hole loss by transport through the oxide. 1 Although the discussion thus far has assumed that the holes (after an initial displacement) are essentially immobile near 80 K, hole motion is greatly accelerated at higher fields. 4 The curves in figure 4 show evidence that the rate of ΔV_{FB} decay is bias dependent: the shift with 10-V applied, shown in curve B (four pulses of 150 krads each) decays more rapidly than the corresponding 5-V shift (curve C); the 20-V shift for one 150-krad pulse (curve D) although initially smaller, decays even more rapidly. If the decay were due to electron injection, the samples biased at 20 V which received multiple pulses should show much more rapid charge loss since the space-charge contribution to the interface field is greater; however, this was not observed (except in the sample that received five 175-krad pulses (curve F), which will be discussed). The charge-loss mechanism here is therefore believed to be field-activated hole transport at the moderately high $(2.3 \times 10^6 \text{ V/cm})$ field present across the oxide.

The sample exposed to five 175-krad pulses at 20-V bias (fig. 4, curve F) shows relatively rapid charge loss which slows down in about a minute. This sample was exposed to the radiation pulses while mounted in an evacuated sample holder maintained at 80 K. The only mechanisms for removing heat from the sample were radiation to the 80-K surroundings (negligible at low temperature) and direct thermal conduction along the two header lead wires to the socket. It is therefore likely that this sample underwent essentially lossless heating when subjected to 875 krads(SiO_2) in 67 ms. The temperature rise was estimated to be ~ 60 K from the low-temperature heat capacity of the

¹H. E. Boesch, Jr., F. B. McLean, J. M. McGarrity, and G. A. Ausman, Jr., IEEE Trans. Nucl. Sci. NS-22, 2163 (1975).

⁴F. B. McLean, H. E. Boesch, Jr., and J. M. McGarrity, Hole Transport and Recovery Characteristics of MOS Gate Insulators, IEEE Nuclear and Space Radiation Effects Conference, July 1976.

silicon substrate and the header material. At $\sim\!140$ K, hole transport is greatly accelerated compared to its rate at 80 K. While this mechanism does not strictly apply to devices operated at 80 K, the possibility that devices may undergo significant transient heating in a real radiation environment should not be ignored.

4.2.2.4 Collapse of Applied Field

Another mechanism that may limit oxide-charge buildup in devices operated in pulsed nuclear environments is collapse of the bias voltage applied across the oxide during a radiation pulse. Since the radiation-generated electrons are mobile, their motion through the oxide under the applied field constitutes a current which may discharge the gate-substrate capacitance, C. If the circuit that consists of the internal resistances of the device and any series external elements has an equivalent series resistance R such that RC is greater than the duration of the radiation pulse τ , the bias on the device will collapse during a radiation pulse of sufficient magnitude. For RC > τ , the fractional charge loss on C due to electron movement only is given by

$$\Delta Q/Q = 1/2\rho AL/CV_G = \rho L^2/2\epsilon V_G$$
 (20)

where A is the oxide area and Q is the charge on the capacitor. The dose D at which complete field collapse (discharge, corresponding to $\Delta Q/Q \sim 1$) should first occur, may be obtained through application of equation (4):

$$D_{\text{sat}} \sim 6 \times 10^{-7} \text{ V}_{G}/\text{f(E)L}^{2}$$
 (21)

The corresponding saturation flatband shift is equal to the magnitude of the applied voltage:

$$|\Delta V_{FB}(D_{sat})| \sim |-V_{G}|$$
 (22)

4.2.2.5 Dielectric Breakdown

Another mechanism may operate if the ones discussed so far fail to limit charge buildup in an MOS structure or if weaknesses exist in that structure. During the course of the experiments, a few MOS capacitors short-circuited under test, usually after receiving a high dose (greater than 3 \times 10 5 rads(SiO2)) and following irradiation at $V_{\rm G} \geq 20$ V. Yang, Johnson, and Lampert 15 have investigated a self-enhanced breakdown mechanism in which electrons tunneling into the SiO2 in high-field regions become "hot" and generate holes by impact ionization, thereby enhancing the local field and therefore the tunneling current. A sudden increase in interface field due to holes generated in the oxide by a radiation pulse may trigger this failure mechanism in an MOS structure with oxide defects (near-pinholes) which cause local field concentration. In complex devices such as LSI logic and CCD's, the device geometry (e.g., metallization pattern) may cause field concentration in small areas, leading to device burnout.

4.2.2.6 Applicability to Higher Temperatures

It should be noted that most of the flatband-shift limitation processes which have been discussed are not inherently restricted to situations in which the device is irradiated at low temperature. With the exception of the hole-transport processes, the modeling and conclusions presented are valid for any situation in which little hole motion takes place during the radiation pulse. The data of Hughes et al³ and Srour et al¹⁰ indicate that little hole motion occurs in less than 10⁻⁸ s even at room temperature. Therefore, most of the processes discussed should limit the initial shift for room temperature irradiations in most pulsed nuclear and flash x-ray environments. At room temperature, rapid charge removal by hole transport ("short-term annealing") takes place within seconds after irradiation. The flatband or threshold shift remaining after transport is presumably determined by the highly processing-dependent density of deep-hole traps in the oxide which capture some of the transported holes. Consequently, any process which limits the initial shift should cause a proportionate reduction in the shift remaining at late time.

¹H. E. Boesch, Jr., F. B. McLean, J. M. McGarrity, and G. A.
Ausman, Jr., IEEE Trans. Nucl. Sci. NS-22, 2163 (1975).

³R. C. Hughes, E. P. Eer Nisse, and H. J. Stein, IEEE Trans. Nucl. Sci. NS-22, 2227 (1975).

¹⁰J. R. Srour, O. L. Curtis, Jr., and K. Y. Chiu, IEEE Trans. Nucl. Sci. NS-21, No. 6, 73 (1974).

¹⁵w. C. Johnson, IEEE Trans. Nucl. Sci. NS-22, 2144 (1975).

5. CONCLUSIONS

The yield of holes produced in $\rm SiO_2$ subjected to high-energy short-pulse radiation at ~ 80 K was measured as a function of electric field, and a hole-electron pair-creation energy of 18 eV was derived. Early differences in $\Delta V_{\rm FB}$ under positive and negative biases at 80 K imply an initial hole displacement of ~ 9 nm at 10^6 V/cm. The extremely large flatband-voltage shifts attained in MOS structures per unit dose at 80 K were shown to be predictable, processing independent, and linear in dose to $\sim 10^5$ rads($\rm SiO_2$). At higher doses the shifts increase sublinearly with dose, primarily as a result of recombination in a low-field region of the oxide. Other mechanisms such as electron injection and hole transport probably play a role as radiation-induced space-charge fields in the oxide become large. The limitation processes should be effective even at room temperature so long as little hole transport takes place during irradiation.

ACKNOWLEDGEMENTS

The authors thank R. Severance and E. Ross of the AFRRI LINAC staff for their beyond-the-call efforts during the experiment and F. B. McLean and G. A. Ausman, Jr., for many helpful discussions. The technical assistance of K. C. Hall is gratefully acknowledged.

LITERATURE CITED

- (1) H. E. Boesch, Jr., F. B. McLean, J. M. McGarrity, and G. A. Ausman, Jr., IEEE Trans. Nucl. Sci. NS-22, 2163 (1975).
- (2) H. H. Sander and B. L. Gregory, IEEE Trans. Nucl. Sci. NS-22, 2157 (1975).
- (3) R. C. Hughes, E. P. Eer Nisse, and H. J. Stein, IEEE Trans. Nucl. Sci. NS-22, 2227 (1975).
- (4) F. B. McLean, H. E. Boesch, Jr., and J. M. McGarrity, Hole Transport and Recovery Characteristics of MOS Gate Insulators, IEEE Nuclear and Space Radiation Effects Conference (July 1976).
- (5) G. A. Ausman, Jr., and F. B. McLean, Appl. Phys. Lett. <u>26</u>, 173 (1975).
- (6) H. E. Boesch, Jr., Development of Apparatus for Performing Rapid Capacitance-Voltage Measurements on MIS Structures, Harry Diamond Laboratories TM-76-33 (December 1976).

LITERATURE CITED (Cont'd)

- (7) F. B. McLean, H. E. Boesch, Jr., P. S. Winokur, J. M. McGarrity, and R. B. Oswald, Jr., IEEE Trans. Nucl. Sci. NS-21, 47 (1974).
- (8) O. L. Curtis, Jr., J. R. Srour, and K. Y. Chiu, J. Appl. Phys. 45, 4506 (1974).
- (9) R. E. Leadon, D. P. Snowden, and J. M. Wilkenfeld Radiation Effects in Semiconductor and Insulator Materials, IRT Corporation, Harry Diamond Laboratories CR-76-152-1 (April 1976).
- (10) J. R. Srour, O. L. Curtis, Jr., and K. Y. Chiu, IEEE Trans. Nucl. Sci. NS-21, No. 6, 73 (1974).
- (11) R. L. Nielsen and D. K. Nichols, IEEE Trans. Nucl. Sci. NS-20, No. 6, 319 (1973).
- (12) E. Harari, S. Wang, and B. S. H. Royce, J. Appl. Phys. <u>46</u>, 1310 (1975).
- (13) J. P. Mitchell, IEEE Trans. Elec. Dev. <u>ED-14</u>, No. 11, 764 (1967).
- (14) R. J. Powell, IEEE Trans. Nucl. Sci. NS-22, 2240 (1975).
- (15) W. C. Johnson, IEEE Trans. Nucl. Sci. NS-22, 2144 (1975).

DISTRIBUTION

DEFENSE DOCUMENTATION CENTER CAMERON STATION, BUILDING 5 ALEXANDRIA, VA 22314 ATTN DDC-TCA (12 COPIES)

COMMANDER
USA RSCH & STD GP (EUR)
BOX 65
FPO NEW YORK 09510
ATTN LTC JAMES M. KENNEDY, JR.
CHIEF, PHYSICS & MATH BRANCH

COMMANDER
US ARMY MATERIEL DEVELOPMENT
5 READINESS COMMAND
5001 EISENHOWER AVENUE
ALEXANDRIA, VA 22333
ATTN DRXAM-TL, HQ TECH LIBRARY
ATTN DRCDE-B, DIR FOR DEV 5 ENGR
ATTN DRCDE-R, SYS EVAL 5 TESTING
ATTN DRCDE-D, LAWRENCE FLYNN

COMMANDER
USA ARMAMENT COMMAND
ROCK ISLAND, IL 61201
ATTN DRSAR-ASF, FUZE DIV
ATTN DRSAR-ROF, SYS DEV DIV - FUZES
ATTN DRSAR-SC, CHIEF SCIENTIST

COMMANDER
USA MISSILE & MUNITIONS CENTER
& SCHOOL
REDSTONE ARSENAL, AL 35809
ATTN ATSK-CTD-F

ARGONNE NATIONAL LABORATORY 9700 SOUTH CASS AVE ARGONNE, IL 60439

BROOKHAVEN NATIONAL LABORATORY ASSOCIATED UNIVERSITIES, INC. UPTON, LONG ISLAND, NY 11973

US DEPARTMENT OF COMMERCE ASSISTANT SECRETARY FOR SCIENCE 6 TECHNOLOGY WASHINGTON, DC 20230

US ENERGY RESEARCH & DEVELOPMENT ADMINISTRATION WASHINGTON, DC 20545 ATTN ASST ADMIN FOR NUCLEAR ENERGY ATTN DIV OF SPACE NUCLEAR SYSTEMS ATTN DIV OF REACTOR RES & DEV

US ENERGY RESEARCH & DEVELOPMENT ADMINISTRATION ALBUQUERQUE OPERATIONS P.O. BOX 5400 ALBUQUERQUE, NM 87115 ATTN DOCUMENT CONTROL FOR WSSB

US ENERGY RESEARCH & DEVELOPMENT ADMINISTRATION TECHNICAL INFORMATION ORGANIZATION P.O. BOX 62 OAK RIDGE, TN 37830

DIRECTOR
ARMED FORCES RADIOBIOLOGY RESEARCH
INSTITUTE
DEFENSE NUCLEAR AGENCY
NATIONAL NAVAL MEDICAL CENTER
BETHESDA, MD 20014
ATTN R. WEISS, CPT.

DIRECTOR
DEFENSE ADVANCED RESEARCH
PROJECTS AGENCY
ARCHITECT BLDG
1400 WILSON BLVD
ARLINGTON, VA 22209
ATTN DIR, MATERIAL SCIENCES
ATTN TECH INFO OFFICE
ATTN DIR, STRATEGIC TECHNOLOGY OFFICE
ATTN DIR, TECHNOLOGY
ASSESSMENTS OFFICE

DEFENSE COMMUNICATION ENGINEER CENTER 1860 WIEHLE AVENUE RESTON, VA 22090 ATTN CODE R320, C. W. BERGMAN ATTN CODE R410, JAMES W. MCLEAN ATTN RES & DEV

DIRECTOR
DEFENSE COMMUNICATIONS AGENCY
WASHINGTON, DC 20305
ATTN CODE 930, MONTE I. BURGETT, JR
ATTN CODE 540.5
ATTN TECH LIBRARY

DIRECTOR
DEFENSE INTELLIGENCE AGENCY
WASHINGTON, DC 20301
ATTN DS-4A2

DIRECTOR

DEFENSE NUCLEAR AGENCY
WASHINGTON, DC 20305
ATTN DDST
ATTN RAEV (3 COPIES)
ATTN STIL TECH LIBRARY
ATTN STVL
ATTN RATN
ATTN PETER HAAS, DEP DIR,
SCI TECHNOLOGY
ATTN AEROSPACE SYS DIR (SPAS)
ATTN VULNERABILITY DIRECTORATE

DIRECTOR
DEFENSE COMMUNICATIONS AGENCY
NATIONAL MILITARY COMMAND
SYSTEM SUPPORT CENTER
WASHINGTON, DC 20301
ATTN TECHNICAL DIRECTOR (B102)

DIR OF DEFENSE RSCH & ENGINEERING
DEPARTMENT OF DEFENSE
WASHINGTON, DC 20301
ATTN DD/S&SS
ATTN DD/S&AS
ATTN AD/E&FS
ATTN DEP DIR (TEST & EVALUATION)
ATTN DEP DIR (RES & ADVANCED TECH)

COMMANDER
FIELD COMMAND
DEFENSE NUCLEAR AGENCY
KIRTLAND AFB, NM 87115
ATTN FCPR

DIRECTOR
INTERSERVICE NUCLEAR WEAPONS SCHOOL
KIRTLAND AFB, NM 87115
ATTN DOCUMENT CONTROL

DIRECTOR
JOINT STRATEGIC TARGET PLANNING STAFF,
JCS
OFFUTT AFB
OMAHA, NB 68113
ATTN JLTW-2

CHIEF
LIVERMORE DIVISION, FIELD COMMAND DNA
LAWRENCE LIVERMORE LABORATORY
F.O. BOX 808
LIVERMORE, CA 94550
ATTN FCPRL

DIRECTOR NATIONAL SECURITY AGENCY FT. GEORGE G. MEADE, MD 20755 ATTN O. O. VAN GUNTEN, R~425 ATTN TDL

OFFICE, CHIEF OF RESEARCH, DEVELOPMENT & ACQUISITION
DEPARTMENT OF THE ARMY
WASHINGTON, DC 20310
ATTN DAMA-AR, CHIEF SCI, DA & DIR OF
ARMY RES, DR. M. E. LASSER
ATTN DAMA-AR, RESEARCH PROGRAMS
ATTN DAMA-RA, SYS REVIEW & ANALYSIS
ATTN DAMA-CSS-D, R&D TEAM

BALLISTIC MISSILE DEFENSE PROGRAM MANAGER OFFICE COMMONWEALTH BUILDING 1300 WILSON BLVD ARLINGTON, VA 22209 ATTN DACS-BMT, TECHNOLOGY DIR

PROJECT MANAGER
ARMY TACTICAL DATA SYSTEMS
US ARMY ELECTRONICS COMMAND
FORT MONMOUTH, NJ 07703
ATTN DWAINE B. HUEWE
ATTN DRCPN-TDS-SD

COMMANDER
BMD SYSTEM COMMAND
PO BOX 1500
HUNTSVILLE, AL 35807
ATTN BDMSC-TEN, NOAH J. HURST
ATTN BMDSC-T, TEST & SYS ENGR SUP DIR

COMMANDER
BALLISTIC MISSILE DEFENSE ADVANCED
TECHNOLOGY CENTER
PO BOX 1500
HUNTSVILLE, AL 35807
ATTN DIRECTOR, ATC-X
ATTN TECH LIB

COMMANDER
US ARMY COMMUNICATIONS COMMAND
FORT HUACHUCA, AZ 85613
ATTN TECH LIB

COMMANDER
US ARMY COMPUTER SYS COMMAND
FORT BELVOIR, VA 22060
ATTN TECH LIB

COMMANDER
US ARMY ELECTRONICS COMMAND
FORT HOLABIRD, MD 21219

PROJECT MANAGER, SAM-D US ARMY MATERIEL DEV & READINESS COMMAND REDSTONE ARSENAL, AL 35809

US ARMY MISSILE RESEARCH, DEV, &
ENGINEERING LABORATORY
US ARMY MISSILE COMMAND
REDSTONE ARSENAL, AL 35809
ATTN DRSMI-RF, ADV SYS CONCEPTS OFC
ATTN DRSMI-RG, GUIDANCE & CONTROL
ATTN DRSMI-RE, ADVANCED SENSORS DIR

The second secon

COMMANDER FRANKFORD ARSENAL BRIDGE AND TACONY STREETS PHILADELPHIA, PA 19137 ATTN SARFA-FCD, MARVIN ELNICK ATTN A1000, TECH DIR

COMMANDING OFFICER NIGHT VISION LABORATORY U. S. ARMY ELECTRONICS COMMAND FORT BELVOIR, VA 22060 ATTN CAPT ALLAN S. PARKER ATTN TECHNICAL LIBRARY

COMMANDER PICATINNY ARSENAL DOVER, NJ 07801 ATTN SARPA-ND-D-B, EDWARD J. ARBER ATTN SARPA-ND-W ATTN SARPA-ND-N ATTN SARPA-FR-E, LOUIS AVRAMI ATTN SARPA-FR-S-P ATTN SARPA-TS-S, SCI & TECH INFO DIV ATTN SARPA-ND-N-E

REDSTONE SCIENTIFIC INFORMATION CTR US ARMY MISSILE COMMAND REDSTONE ARSENAL, AL 35809 ATTN CHIEF, DOCUMENTS

SECRETARY OF THE ARMY WASHINGTON, DC 20310 ATTN ODUSA OR DANIEL WILLIARD

ASSISTANT SECRETARY OF THE ARMY (R&D) WASHINGTON, DC 20310 ATTN DEP FOR SCI & TECH

TRASANA WHITE SANDS MISSILE RANGE, NM 88002 ATTN ATAA-EAC, FRANCIS N. WINANS

US ARMY BALLISTIC RESEARCH LABORATORIES ABERDEEN PROVING GROUND, MD 21005 ATTN DAVID L. RIGOTTI, DRXBR-BVL ATTN DRXBR-X, JULIUS J. MESZAROS ATTN DRXBR-AM, W. R. VANANTWERP ATTN DRXBR-VL, JOHN W. KINCH ATTN DRXBR-VL, ROBERT L. HARRISON

US ARMY COMMUNICATIONS SYSTEMS AGENCY FORT MONMOUTH, NJ 07703 ATTN SCCM-AD-SV (LIBRARY)

COMMANDER US ARMY ELECTRONICS COMMAND FORT MONMOUTH, NJ 07703 ATTN DRSEL-CT-HDK, ABRAHAM E. COHEN ATTN DRSEL-TL-MD, GERHART K. GAULE ATTN DRSEL-GG-TD, W. R. WERK ATTN DRSEL-PL-ENV, HANS A. BOMKE ATTN DRSEL-TL-ND, S. KRONENBEY ATTN DRSEL-TL-IR, EDWIN T. HUNTER ATTN DRSEL-TL-EN, ROBERT LUX ATTN DRSEL-CE, T. PREIFFER ATTH DRSEL-RD, DIR, RES, DEV, 6 ENGR ATTH DRSEL-WL-D, ELECTP WARFARE LAB ATTH DRSEL-VL-D, AVIONICS LAB ATTH DRSEL-TL-D, ELECTRONICS TECHNOLOGY 6 DEVICES LAB COMMANDANT US ARMY ENGINEER SCHOOL FT. BELVOIR, VA 22060

COMMANDER-IN-CHIEF US ARMY EUROPE AND SEVENTH ARMY APO NEW YORK 09403 ATTN ODCSE-E, AEAGE-PI

COMMANDANT U.S. ARMY FIELD ARTILLERY SCHOOL FORT SILL, OK 73503 ATTN ATSFA-CTD-ME, HARLEY MOBERG

COMMANDER US ARMY MISSILE COMMAND REDSTONE ARSENAL HUNTSVILLE, AL 35809 ATTN DRCPM-PE-EA, WALLACE O. WAGNER ATTN DRSI-RGP, VICTOR W. RUWE ATTN DRSMI-RRR, FAISON P. GIBSON ATTN DRCPM-MDTI, CPT JOE A SIMS ATTN DRCPM-LCEX, HOWARD H. HENRIKSEN ATTN TECHNICAL LIBRARY ATTN DRSMI-RGP, HUGH GREEN ATTN ARMY MISSILE RDE LAB ATTN TECH LIB

COMMANDER US ARMY MOBILITY EQUIP R&D CTR FORT BELVOIR, VA 22060 ATTN STSFB-MW, JOHN W. BOND, JR.

US ARMY NUC AND CHEMICAL SURETY GP BLDG, 2073, NORTH AREA FT. BELVOIR, VA 22060 ATTN MOSG-ND, MAJ SIDNEY W. WINSLOW

COMMANDER US ARMY NUCLEAR AGENCY FORT BLISS, TX 79916 ATTN ATCN-W, LTC LEONARD A. SLUGA

COMMANDER U.S. ARMY TANK AUTOMOTIVE COMMAND WARREN, MI 48090 ATTN DRCPM-GCM-SW, LYLE A. WOLCOTT

US ARMY TEST AND EVALUATION COMMAND ABERDEEN PROVING GROUND, MD 21005 ATTN DRSTE-EL, R. I. KOLCHIN ATTN DRSTE-NB, R. R. GALASSO

COMMANDER WHITE SANDS MISSILE RANGE WHITE SANDS MISSILE RANGE, NM 88002 ATTN STEWS-TE-NT. MR. MARVIN P. SQUIRES

COMMANDER COMMANDER
US ARMY NUCLEAR AGENCY
FORT BLISS, TX 79916
ATTN ATCN-W, WEAPONS EFFECTS DIV
ATTN TECH LIB

COMMANDER NAVAL RESEARCH LABORATORY HO, US ARMY TEST & EVALUATION COMMAND WASHINGTON, DC 20375 ABERDEEN PROVING GROUND, MD 21005 ATTN DRSTE-EL, ELECTRONICS MATERIEL TESTING DIR

COMMANDER EDGEWOOD ARSENAL EDGEWOOD ARSENAL, MD 21010 ATTN SAREA-R, RES LABORATORIES ATTN SAREA-DE, DIR OF DEV & ENGR ATTN SAREA-TS, TECH LIB

OFFICE OF RESEARCH, DEVELOPMENT, TEST & EVALUATION DEPT OF THE NAVY WASHINGTON, DC 20360 ATTN OP-987, R&D PLANS

US NAVAL ACADEMY ENGINEERING DEPT ANNAPOLIS, MD 21402 ATTN LIBRARY

COMMANDER NAVAL AIR SYSTEMS COMMAND HQ DEPT OF THE NAVY WASHINGTON, DC 20360

COMMANDER NAVAL ELECTRONICS LABORATORY CENTER SAN DIEGO, CA 92152 ATTN CODE 6400, TECH INFO DIV

SUPERINTENDANT NAVAL POSTGRADUATE SCHOOL MONTEREY, CA 93940 ATTN LIBRARY, CODE 2124

CHIEF OF NAVAL RESEARCH NAVY DEPARTMENT ARLINGTON, VA 22217 ATTN CODE 427 ATTN CODE 421, DORAN W. PADGETT ATTN TECHNICAL LIBRARY

COMMANDING OFFICER NAVAL AVIONICS FACILITY 21ST AND ARLINGTON AVENUE INDIANAPOLIS, IN 46218 ATTN BRANCH 942, D. J. REPASS

COMMANDER NAVAL ELECTRONIC SYSTEMS COMMAND HEADQUARTERS WASHINGTON, DC 20360 ATTN PME 117-21 ATTN ELEX 05323, CLEVELAND F. WATKINS ATTN CODE 504510 ATTN CODE 5032, CHARLES W. NEILL ATTN CODE 504511, CHARLES R. SUMAN ATTN CODE 9053, TECH INFO SVCS BR

COMMANDING OFFICER NAVAL INTELLIGENCE SUPPORT CENTER 4301 SUITLAND ROAD, BLDG 5 WASHINGTON, DC 20390 ATTN P. ALEXANDER ATTN NISC-45

DIRECTOR ATTN 6601, E. WOLICKI ATTN CODE 6631, JAMES C. RITTER ATTN CODE 4004, EMANUAL L. BRANCATO ATTN CODE 2627, DORIS R. FOLEN ATTN CODE 7701, JACK D. BROWN ATTN CODE 5216, HAROLD L. HUGHES ATTN CODE 5210, JOHN E. DAVEY

NAVAL RESEARCH LAB (Cont'd) ATTN CODE 6440 GEORGE SIGEL ATTN CODE 2620, LIBRARY ATTN CODE 4000, RESEARCH DEPT ATTN CODE 6620, RADIATION EFFECTS

COMMANDER
NAVAL SEA SYSTEMS COMMAND
NAVY DEPARTMENT
WASHINGTON, DC 20362
ATTN SEA-9931, RILEY B. LANE
ATTN SEA-9931, SAMUEL A. BARHAM
ATTN SEA-09632, TECH LIB

NAVAL SHIP ENGINEERING CENTER DEPT OF THE NAVY WASHINGTON, DC 20362 ATTN CODE 6174D2, EDWARD F. DUFFY

COMMANDER
NAVAL SURFACE WEAPONS CENTER
WHITE OAK, SILVER SPRING, MD 20910
ATTN CODE WX21, TECH LIB
ATTN CODE WA501, NAVY NUC PRGMS OFF
ATTN WA50
ATTN CODE WA52, R. A. SMITH
ATTN CODE WR, RESEARCH &

COMMANDER
NAVAL SURFACE WEAPONS CENTER
DAHLGREN LABORATORY
DAHLGREN, VA 22448
ATTN WILLIAM H. HOLT
ATTN DX-21, LIBRARY DIV

TECHNOLOGY DEPT

COMMANDER
NAVAL WEAPONS CENTER
CHINA LAKE, CA 93555
ATTN CODE 533, TECHNICAL LIBRARY

COMMANDING OFFICER
NAVAL WEAPONS EVALUATION FACILITY
KIRTLAND AIR FORCE BASE
ALBUQUERQUE, NM 87117
ATTN CODE ATG, MR. STANLEY

COMMANDING OFFICER
NAVAL WEAPONS SUPPORT CENTER
CRANE, IN 47522
ATTN CODE 7024, JAMES RAMSEY
ATTN CODE 70242, JOSEPH A. MUNARIN

COMMANDING OFFICER
NUCLEAR WEAPONS TRAINING CENTER PACIFIC
NAVAL AIR STATION, NORTH ISLAND
SAN DIEGO, CA 92135
ATTN CODE 50

DIRECTOR
STRATEGIC SYSTEMS PROJECT OFFICE
NAVY DEPARTMENT
WASHINGTON, DC 20376
ATTN SP2701, JOHN W. PITSENBERGER
ATTN NSP-2342, RICHARD L. COLEMAN
ATTN NSP-27331, PHIL SPECTOR

ASSISTANT SECRETARY OF THE AIR FORCE (RESEARCH & DEVELOPMENT) WASHINGTON, DC 20330

DEPUTY CHIEF OF STAFF, RES & DEV US AIR FORCE WASHINGTON, DC 20330 ATTN RDQPN, S/V & NUCLEAR PROG DIV COMMANDER
AERONAUTICAL SYSTEMS DIC
WRIGHT-PATTERSON AFB, OH 45433

COMMANDER-IN-CHIEF
AEROSPACE DEFENSE COMMAND
ENT AIR FORCE BASE, CO 80912
ATTN TECHNICAL LIBRARY

COMMANDER AEROSPACE RESEARCH LABORATORIES WRIGHT-PATTERSON AFF, OH 45433 ATTN LS, SOLID STATE PHYSICS RES LAB

DIRECTOR

AF AVIONICS LABORATORY

WRIGHT-PATTERSON AFB, OH 45433

ATTN TE, ELECTRONIC TECHNOLOGY DIV

ATTN TER, ELECTRONIC RES BR

ATTN TSR, STINFO BR

ATTN DHE, H. J. HENNECKE

ATTN DHM, C. FRIEND

ATTN DH, LTC MCKENZIE

ATTN DH, LTC MCKENZIE

ATTN ATA, MASON FRIAR

COMMANDER
AF CAMBRIDGE RESEARCH LABORATORIES,
AFSC
L. G. HANSCOM FIELD
BEDFORD, MA 01730
ATTN LQ, SOLID-STATE SCI LAB

COMMANDER
AF FLIGHT DYNAMICS LAB
WRIGHT-PATTERSON AFF, OH 45433
ATTN PTS, SURVIVABILITY/
VULNER-BILITY BR
ATTN STS, TECH INFO BR

AF GEOPHYSICS LABORATORY, AFSC HANSCOM AFB, MA 01731 ATTN J. EMERY CORMIER ATTN LGD-STOP 30, FREEMAN SHEPHERD ATTN LCR, EDWARD A. BURKE

AF INSTITUTE OF TECHNOLOGY, AU WRIGHT-PATTERSON AFB, OH 45433 ATTN ENP, CHARLES J. BRIDGMAN

AF MATERIALS LABOTATORY, AFSC WRIGHT-PATTERSON AFB, OH 45433 ATTN LTE

KIRTLAND AFB, NM 87117
ATTN ELA
ATTN SAB
ATTN ELP, TREF SECTION
ATTN ELP, JULIAN NICHOLS
ATTN DEX
ATTN ELS
ATTN NTS
ATTN ELXT
ATTN SELXT
ATTN SEL, NUCLEAR SYS DIV
ATTN ELC, COMM & SATELLITE BR

AF WEAPONS LABORATORY, AFSC

AFTAC PATRICK AFB, FL 32925 ATTN TFS, MAJ MARION F. SCHNEIDER ATTN TAE COMMANDER
ASD
WPAFE, OH 45433
ATTN ASD/ENESS, PETER T. MARTH
ATTN ASD-YH-EX, LTC ROBERT LEVERETTE
ATTN ENACC, ROBERT L, FISH

HEADQUARTERS ELECTRONIC SYSTEMS DIVISION, (AFSC) HANSCOM AFB, MA 01731 ATTN YWEI ATTN YSEV

COMMANDER FOREIGN TECHNOLOGY DIVISION, AFSC WRIGHT-PATTERSON AFB, OH 45433 ATTN ETET, CAPT RICHARD C. HUSEMANN

COMMANDER
ROME AIR DEVELOPMENT CENTER, AFSC
GRIFFISS AFB, NY 13440
ATTN RBRAC, I. L. KRULAC
ATTN RBRP, CLYDE LANE
ATTN TUT, TEST & EVAL BR
ATTN TUT, TEST & EVAL BR
ATTN TUM, MATERIEL BR
ATTN TIL, TECHNICAL LIBRARY

COMMANDER
ROME AIR DEVELOPMENT CENTER, AFSC
HANSCOM AFB, MA 01731
ATTN ET, R. BUCHANAN
ATTN ETS, R. DOLAN
ATTN ETS, A KAHAN

DIRECTOR
AF OFFICE OF SCIENTITIC RESEARCH
1400 WILSON BLVD
ARLINGTON, VA 22209
ATTN NE, DIR OF ELECTRONIC
5 SOLID STATE SCI

COMMANDER
SPACE & MISSILE SYSTEMS ORGANIZATION
(SAMSO)
PO BOX 92960
WORLDWAY POSTAL CENTER
LOS ANGELES, CA 90009
ATTN RS, DEP FOR REENTRY SYS
ATTN LV, DEP FOR LAUNCH VEHICLES
ATTN SK, DEP FOR COMM SYS

COMMANDER
AF SPECIAL WEAPONS CENTER (OAS)
KIRTLAND AFB, NM 87117
ATTN TE, TEST & EVAL SYS PROG OFC

COMMANDER
HQ AIR FORCE SYSTEMS COMMAND
ANDREWS AFB
WASHINGTON, DC 20331
ATTN TECHNICAL LIBRARY

SAMSO/DY
POST OFFICE BOX 92960
WORLDWAY POSTAL CENTER
LOS ANGELES, CA 90009
ATTN DYS, MAJ LARRY A. DARDA
ATTN DYS, CAPT WAYNE SCHOBER
ATTN DYS, CAPT E. MERZ

SAMSO/IN
POST OFFICE BOX 92960
WORLDWAY POSTAL CENTER
LOS ANGELES, CA 90009
ATTN IND, I. J. JUDY

SAMSO/MN NORTON AFB, CA 92409 ATTN MNNG, CAPT DAVID J. STROBEL ATTN MNNH

SAMSO/RS
POST OFFICE BOX 92960
WORLDWAY POSTAL CENTER
LOS ANGELES, CA 90009
ATTN RSSE, LTC KENNETH L. GILBERT
ATTN RSMG, CAPT COLLIER

SAMSO/SK POST OFFICE BOX 92960 WORLDWAY POSTAL CENTER LOS ANGELES, CA 90009 ATTN SKF, PETER J. STADLER

SAMSO/SZ POST OFFICE BOX 92960 WORLDWAY POSTAL CENTER LOS ANGELES, CA 90009 ATTN SZJ, CAPT JOHN H. SALCH

COMMANDER IN CHIEF STRATEGIC AIR COMMAND OFFUTT AFB, NB 68113 ATTN NRI-STINFO LIBRARY ATTN XPFS, MAJ BRIAN STEPHAN

UNIVERSITY OF CALIFORNIA
LANRENCE LIVERMORE LABORATORY
PO BOX 808
LIVERMORE, CA 94550
ATTN DONALD J. MEEKER, L-545
ATTN JOSEPH E. KELLER, JR., L-125
ATTN RONALD L. OTT, L-531
ATTN HANS KRUGER, L-96
ATTN LAWRENCE CLELAND, L-156
ATTN FREDERICK R. KOVAR, L-31
ATTN TECH INFO DEPT, L-3

LOS ALAMOS SCIENTIFIC LABORATORY P.O. BOX 1663 LOS ALAMOS, NM 87544 ATTN MARVIN M. HOFFMAN ATTN J. ARTHUR FREED ATTN BRUCE W. NOEL

SANDIA LABORATORIES LIVERMORE LABORATORY PO BOX 969 LIVERMORE, CA 94550 ATTN THEODORE A. DELLIN

SANDIA LABORATORIES
PO BOX 5800
ALBUQUERQUE, NM 87115
ATTN 3141 SANDIA RPT COLL
ATTN ORG 2110, J. A. HOOD
ATTN JACK V. WALKER, 5220
ATTN ORG 1933, F. N. COPPAGE
ATTN DIV 5231, JAMES H. RENKEN
ATTN R. GREGORY, ORG 2140

ADMINISTRATOR
NASA HEADQUARTERS
WASHINGTON, DC 20546
ATTN OFC OF AERONAUTICS
AND SPACE TECHNOLOGY

AMES RESEARCH CENTER NASA MOFFETT FIELD, CA 94035 ATTN DIR OF RESEARCH SUPPORT DIRECTOR
NASA
GODDARD SPACE FLIGHT CENTER
GREENBELT, MD 20771
ATTN 250, TECH INFO DIV

DIRECTOR NSAS HUGH L. DRYDEN FLIGHT RESEARCH CENTER EDWARDS, CA 93523 ATTN TECHNICAL LIBRARY

DIRECTOR
NASA
JOHN F. KENNEDY SPACE CENTER, FL 32899
ATTN TECHNICAL LIBRARY

DIRECTOR
NASA
LEWIS RESEARCH CENTER
CLEVELAND, OH 44135
ATTN TECHNICAL LIBRARY

JET PROPULSION LABORATORY CALIFORNIA INSTITUTE OF TECHNOLOGY 4800 OAK GROVE DRIVE PASADENA, CA 91103 ATTN TECHNICAL LIBRARY

DIRECTOR
NASA
LANGLEY RESEARCH CENTER
HAMPTON, VA 23665
ATTN TECHNICAL LIBRARY
ATTN INSTR RES DIV

DIRECTOR
NASA
GEORGE C. MARSHALL SPACE FLIGHT CENTER
MARSHALL SPACE FLIGHT CENTER, AL 35812
ATTN EA, SCI & ENGR
ATTN EC, ELECTRONICS & CONTROL
ATTN EC-21, GUIDANCE, CONTROL
6 INSTR DIV
ATTN EC-41, ELECTRONICS DEV DIV
ATTN ES-21, RADIATION & LOW TEMP SCI
ATTN ES-31, PHYSICS & INSTR DIV

CENTRAL INTELLIGENCE AGENCY ATTN: RD/SI, RM 5G48, HQ BLDG WASHINGTON, DC 20505 ATTN ALICE A. PADGETT

DEPARTMENT OF COMMERCE
NATIONAL BUREAU OF STANDARDS
WASHINGTON, DC 20234
ATTN APPL RAD DIV, ROBERT C. PLACIOUS
ATTN JUDSON C. FRENCH

AEROJET ELECTRO-SYSTEMS CO. DIV. AEROJET-GENERAL CORPORATION P.O. BOX 296 AZUSA, CA 91702 ATTN THOMAS D. HANSCOME, B170/D6711

AEROSPACE CORPORATION
PO BOX 92957
LOS ANGELES, CA 90009
ATTN IRVING M. GARFUNKEL
ATTN JULIAN REINHEIMER
ATTN LIBRARY
ATTN MELVIN J. BERNSTEIN
ATTN WILLIAM W. WILLIS
ATTM S. P. BOWER
ATTN JOHN DITRE
ATTN L. W. AUKERMAN

ANALOG TECHNOLOGY CORPORATION 3410 EAST POOTHILL BOULEVARD PASADENA, CA 91107 ATTN JOHN JOSEPH BAUM

AVCO RESEARCH & SYSTEMS GROUP 201 LOWELL STREET WILMINGTON, MA 01887 ATTN RESEARCH LIB, A830, RM 7201

BDM CORPORATION, THE 1920 ALINE AVE VIENNA, VA 22180 ATTN T. H. NEIGHBORS

BDM CORPORATION, THE PO BOX 9274 ALBUQUERQUE INTERNATIONAL ALBUQUERQUE, NM 87119 ATTN D. R. ALEXANDER

BENDIX CORPORATION, THE COMMUNICATION DIVISION EAST JOPPA ROAD - TOWSON BALTIMORE, MD 21204 ATTN DOCUMENT CONTROL

BENDIX CORPORATION, THE RESEARCH LABORATORIES DIV BENDIX CENTER SOUTHFIELD, MI 48075 ATTN MGR PRGM DEV, DONALD J. NIEHAUS ATTN MAX FRANK

BOEING COMPANY, THE PO BOX 3707 SEATTLE, WA 98124 ATTN HOWARD W. WICKLEIN, MS 17-11 ATTN DAVID DYE, MS 87-75 ATTN AEROSPACE LIBRARY ATTN ROBERT S. CALDWELL, 2R-00

BOOZ-ALLEN AND HAMILTON, INC. 106 APPLE STREET NEW SHREWSBURY, NJ 07724 ATTN R. J. CHRISNER

CALIFORNIA INSTITUTE OF TECHNOLOGY JET PRCPULSION LABORATORY 4800 OAK PARK GROVE PASADENA, CA 91103 ATTN J. BRYDEN ATTN A. G. STANLEY

CHARLES STARK DRAPER LABORATORY INC. 68 ALBANY STREET CAMBRIDGE, MA 02139 ATTN KENNETH FERTIG ATTN PAUL R. KELLY ATTN RICHARD G. HALTMAIER

CINCINNATI ELECTRONICS CORPORATION 2630 GLENDALE - MILFORD ROAD CINCINNATI, OH 45241 ATTN C. R. STUMP ATTN LOIS HAMMOND

COMPUTER SCIENCES CORPORATION 201 LA VETA DRIVE, NE ALBUQUERQUE, NM 87108 ATTN RICHARD H. DICKHAUT

CUTLER-HAMMER, INC.
AIL DIVISION
COMAC ROAD
DEER PARK, NY 11729
ATTN CENTRAL TECH FILES, ANN ANTHONY

DIKEWOOD CORPORATION, THE 1009 BRADBURY DRIVE, SE UNIVERSITY RESEARCH PARK ALBUQUERQUE, NM 87106 ATTN L. WAYNE DAVIS

E-SYSTEMS, INC. GREENVILLE DIVISION PO BOX 1056 GREENVILLE, TX 75401 ATTN LIBRARY 8-50100

EFFECTS TECHNOLOGY, INC. 5383 HOLLISTER AVENUE SANTA BARBARA, CA 93105 ATTN EDWARD JOHN STEELE

EXPERIMENTAL AND MATHEMATICAL PHYSICS CONSULTANTS P. 0. BOX 66331 LOS ANGELES, CA 90066 ATTN THOMAS M. JORDAN

FAIRCHILD CAMERA AND INSTRUMENT CORPORATION 464 ELLIS STREET MOUNTAIN VIEW, CA 94040 ATTN 2-233, MR. DAVID K. MYERS

FAIRCHILD INDUSTRIES, INC. SHERMAN FAIRCHILD TECHNOLOGY CENTER 20301 CENTURY BOULEVARD GERMANTOWN, MD 20767 ATTN MGR CONFIG DATA & STANDARDS

FLORIDA, UNIVERSITY OF AN INSTITUTION OF EDUCATION ATTH: PATRICIA B. RAMBO P.O. BOX 284 GAIMESVILLE, FL 32601 ATTN D. P. KENNEDY

FORD AEROSPACE & COMMUNICATIONS CORP 3939 FABIAN WAY PALO ALTO, CA 94303 ATTN SAMUEL R. CRAWFORD, MS 531 ATTN EDWARD R. HAHN, MS-X22 ATTN DONALD R. MCMORROW, MS G30

FORD AEROSPACE & COMMUNICATIONS OPERATIONS FORD & JAMBOREE ROADS NEWPORT BEACH, CA 92663 ATTN KEN C. ATTINGER ATTN E. R. PONCELET, JR ATTN TECH INFO SECTION

FRANKLIN INSTITUTE, THE 20TH STREET AND PARKWAY PHILADELPHIA, PA 19103 ATTN RAMIE H. THOMPSON

GARRETT CORPORATION P.O. BOX 92248 LOS ANGELES, CA 90009 ATTN ROBERT E. WEIR, DEPT. 93-9 GENERAL DYNAMICS CORP
ELECTRONICS DIV ORLANDO OPERATIONS
PO BOX 2566
ORLANDO, FL 32802
ATTN D. W. COLEMAN

GENERAL ELECTRIC COMPANY
SPACE DIVISION
VALLEY FORGE SPACE CENTER
GODDARD BLVD KING OF PRUSSIA
P.O. BOX 8555
PHILADELPHIA, PA 19101
ATTN LARRY I. CHASEN
ATTN JOSEPH C. PEDEN, CCF 8301
ATTN JOHN L. ANDREWS

GENERAL ELECTRIC COMPANY
RE-ENTRY & ENVIRONMENTAL SYSTEMS DIV
PO BOX 7722
3198 CHESTNUT STREET
PHILADELPHIA, PA 19101
ATTN JOHN W. PALCHEFSHY, JR.
ATTN ROBERT V. BENEDICT
ATTN RAY E. ANDERSON

GENERAL ELECTRIC COMPANY ORDNANCE SYSTEMS 100 PLASTICS AVENUE PITTSFIELD, MA 01201 ATTN JOSEPH J. REIDL

GENERAL ELECTRIC COMPANY
TEMPO-CENTER FOR ADVANCED STUDIES
816 STATE STREET (PO DRAWER QQ)
SANTA BARBARA, CA 93102
ATTN DASIAC
ATTN ROYDEN R. RUTHERFORD
ATTN M ESPIG

GENERAL ELECTRIC COMPANY PO BOX 1122 SYRACUSE, NY 13201 ATTN CSP 0-7, L. H. DEE

GENERAL ELECTRIC COMPANY AIRCRAFT ENGINE GROUP EVENDALE PLANT CINCINNATI, OH 45215 ATTN JOHN A. ELLERHORST, E2

GENERAL ELECTRIC COMPANY
AEROSPACE ELECTRONICS SYSTEMS
FRENCH ROAD
UTICA, NY 13503
ATTN CHARLES M. HEWISON, DROP 624
ATTN W. J. PATTERSON, DROP 233

GENERAL ELECTRIC COMPANY PO BOX 5000 BINGHAMTON, NY 13902 ATTN DIVID W. PEPIN, DROP 160

GENERAL ELECTRIC COMPANY-TEMPO ATTN: DASIAC C/O DEFENSE NUCLEAR AGENCY WASHINGTON, DC 20305 ATTN WILLIAM ALFONTE

GENERAL RESEARCH CORPORATION P.O. BOX 3587 SANTA BARBARA, CA 93105 ATTN ROBERT D. HILL GEORGIA INSTITUTE OF TECHNOLOGY GEORGIA TECH RESEARCH INSTITUTE ATLANTA, GA 30332 ATTN R. CURRY

GRUMMAN AEROSPACE CORPORATION SOUTH OYSTER BAY ROAD BETHPAGE, NY 11714 ATTN JERRY ROGERS, DEPT 533

GTE SYLVANIA, INC.
ELECTRONICS SYSTEMS GRP-EASTERN DIV
77 A STREET
NEEDHAM, MA 02194
ATTN CHARLES A. THORNHILL, LIBRARIAN
ATTN LEONARD L. BLAISDELL
ATTN JAMES A. WALDON

GTE SYLVANIA, INC.
189 B STREET
NEEDHAM HEIGHTS, MA 02194
ATTN CHARLES H. RAMSBOTTOM
ATTN HERBERT A. ULLMAN
ATTN H & V GROUP
ATTN PAUL B. FREDRICKSON

GULTON INDUSTRIES, INC. ENGINEERED MAGNETICS DIVISION 13041 CERISE AVENUE HAWTHORNE, CA 90250 ATTN ENGMMAGNETICS DIV

HARRIS CORPORATION
HARRIS SEMICONDUCTOR DIVISION
P.O. BOX 883
MELBOURNE, FL 32901
ATTN C. F. DAVIS, MS 17-220
ATTN T. CLARK, MS 4040
ATTN WAYNE E. ABARE, MS 16-111

HAZELTINE CORPORATION
PULASKI ROAD
GREEN LAWN, NY 11740
ATTN TECH INFO CTR, M. WAITE

HONEYWELL INCORPORATED GOVERNMENT AND AERONAUTICAL PRODUCTS DIVISION 2600 RIDGEWAY PARKWAY MINNEAFOLIS, MN 55413 ATTN RONALD R. JOHNSON, A1622 ATTN R. J. KELL, MS 52572

HONEYWELL INCORPORATED AEROSPACE DIVISION 13350 US HIGHWAY 19 ST. PETERSBURG, FL 33733 ATTN HARRISON H. NOBLE, MS 725-5A ATTN MS 725-J, STACEY H. GRAFF

HONEYWELL INCORPORATED RADIATION CENTER 2 FORBES ROAD LEXINGTON, MA 02173 ATTN TECHNICAL LIBRARY

HUGHES AIRCRAFT COMPANY
CENTINELLA & TEALE
CULVER CITY, CA 90230
ATTN M.S. D157, KEN WALKER
ATTN B. W. CAMPBELL, M.S. 6-E110
ATTN DAN BINDER, MS 6-D147
ATTN JOHN B. SINGLETARY, MS 6-D133

HUGHES AIRCRAFT COMPANY SPACE SYSTEMS DIVISION P.O. BOX 92919 LOS ANGELES, CA 90009 ATTN WILLIAM W. SCOTT, MS A1080 ATTN EDWARD C. SMITH, MS A620

IBM CORPORATION ROUTE 17C OWEGO, NY 13627 ATTN HARRY W. MATHERS, DEPT M41 ATTN FRANK FRANKOVSKY

INTL TEL & TELEGRAPH CORPORATION 500 WASHINGTON AVENUE NUTLEY, NJ 07110 ATTN ALEXANDER T. RICHARDSON

ION PHYSICS CORPORATION SOUTH BEDFORD STREET BURLINGTON, MA 01803 ATTN ROBERT D. EVANS

IRT CORPORATION
P.O. BOX 81087
SAN DIEGO, CA 92138
ATTN LEO D. COTTER
ATTN RALPH H. STAHL
ATTN JAMES A. NABER
ATTN R. L. MERTZ
ATTN MDC

JAYCOR 205 S. WHITING STREET, SUITE 500 ALEXANDRIA, VA 22304 ATTN ROBERT SULLIVAN ATTN CATHERINE TURESKO

JOHNS HOPKINS UNIVERSITY APPLIED PHYSICS LABORATORY JOHNS HOPKINS ROAD LAUREL, MD 20810 ATTN PETER E. PARTRIDGE

KAMAN SCIENCES CORPORATION P.O. BOX 7463 COLORADO SPRINGS, CO 80933 ATTN DONALD H. BRYCE ATTN ALBERT P. BRIDGES ATTN WALTER E. WARE ATTN JOHN R. HOFFMAN ATTN JERRY I. LUBELL

LITTON SYSTEMS, INC.
GUIDANCE & CONTROL SYSTEMS DIVISION
5500 CANOGA AVENUE
WOODLAND HILLS, CA 91364
ATTN VAL J. ASHBY, MS 67
ATTN JOHN P. RETZLER

LITTON SYSTEMS, INC. ELECTRON TUBE DIVISION 1035 WESTMINISTER DRIVE WILLIAMSPORT, PA 17701 ATTN FRANK J. MCCARTHY

LOCKHEED MISSILES AND SPACE
COMPANY, INC.
P.O. BOX 504
SUNNYVALE, CA 94088
ATTN G. F. HEATH, D/81-14
ATTN PHILIP J. HART, DEPT 81-14
ATTN BENJAMIN T. KIMURA, DEPT 81-14
ATTN EDWIN A. SMITH, DEPT 85-85
ATTN L. ROSSI, DEPT 81-64

LOCKHEED MISSILES AND SPACE (Cont'd) ATTN G. H. MORRIS, 81-01 ATTN DEPT 85-85, SAMUEL I. TAIMUTY

LOCKHEED MISSILES AND SPACE COMPANY 3251 HANOVER STREET PALO ALTO, CA 94304 ATTN TECH INFO CTR D/COLL

LTV AEROSPACE CORPORATION VOUGHT SYSTEMS DIVISION P.O. BOX 6267 DALLAS, TX 75222 ATTN TECHNICAL DATA CENTER

LTV AEROSPACE CORPORATION PO BOX 5907 DALLAS, TX 75222 ATTN TECHNICAL DATA CTR

M.I.T. LINCOLN LABORATORY P.O. BOX 73 LEXINGTON, MA 02173 ATTN LEONA LOUGHLIN, LIBRARIAN, A-082

MARTIN MARIETTA AEROSPACE
ORLANDO DIVISION
P.O. BOX 5837
ORLANDO, FL 32805
ATTN MONA C. GRIFFITH, LIB MP-30
ATTN WILLIAM W. MRAS, MP-413
ATTN JACK M. ASHFORD, MP-537

MARTIN MARIETTA CORPORATION
DENVER DIVISION
PO BOX 179
DENVER, CO 80201
ATTN J. E. GOODWIN, MAIL 0452
ATTN RESEARCH LIB, 6617, J. R. MCKEE
ATTN PAUL G. KASE, MAIL 8203
ATTN BEN J. GRAHAM, MS PO-454

MCDONNELL DOUGLAS CORPORATION POST OFFICE BOX 516 ST. LOUIS, MO 63166 ATTN TOM ENDER ATTN TECHNICAL LIBRARY

MCDONNELL DOUGLAS CORPORATION 5301 BOLSA AVENUE HUNTINGTON BEACH, CA 92647 ATTN STANLEY SCHNEIDER

MCDONNELL DOUGLAS CORPORATION 3855 LAKEWOOD BOULEVARD LONG BEACH, CA 90846 ATTN TECHNICAL LIBRARY, C1-290/36-84

MISSION RESEARCH CORPORATION 735 STATE STREET SANTA BARBARA, CA 93101 ATTN WILLIAM C. HART

MISSION RESEARCH CORPORATION-SAN DIEGO
7650 CONVOY COURT
SAN DIEGO, CA 92111
ATTN V. A. J. VAN LINT
ATTN J. P. RAYMOND

RAYTHEON COMPANY
HARTWELL ROAD
BEDFORD, MA 017
ATTN GAJANAN H

MITRE CORPORATION, THE P.O. BOX 208 BEDFORD, MA 01730 ATTN LIBRARY ATTN M. F. FITZGERALD NATIONAL ACADEMY OF SCIENCES 2101 CONSTITUTION AVE, WASHINGTON,DC 20418 ATTN DR. R. S. SHANE, NAT MATERIALS ADVISORY BO

NEW MEXICO, UNIVERSITY OF DEPT OF CAMPUS SECURITY AND POLICE 1821 ROMA NE ALBUQUERQUE, NM 87106 ATTN W. W. GRANNEMANN

NEW MEXICO, UNIVERSITY OF ELECTRICAL ENGINEERING & COMPUTER SCIENCE DEPT ALBUQUERQUE, NM 87131 ATTN HAROLD SOUTHWARD

NORTHROP CORPORATION
ELECTRONIC DIVISION
1 RESEARCH PARK
PALOS VERDES PENINSULA, CA 90274
ATTN JOHN M. REYNOLDS
ATTN VINCENT R. DEMARTINO
ATTN BOYCE T. AHLPORT
ATTN GEORGE H. TOWNER

NORTHROP CORPORATION
NORTHROP RESEARCH AND TECHNOLOGY CENTER
3401 WEST BROADWAY
HAWTHORNE, CA 90250
ATTN DAVID N. POCOCK
ATTN J. R. SROUR
ATTN ORLIE L. CURTIS

NORTHROP CORPORATION ELECTRONIC DIVISION 2301 WEST 120TH STREET HAWTHORNE, CA 90250 ATTN JOSEPH D. RUSSO

PALISADES INST FOR RSCH SERVICES INC. 201 VARICK STREET NEW YORK, NY 10014 ATTN RECORDS SUPERVISOR

PHYSICS INTERNATIONAL COMPANY 27C0 MERCED STREET SAN LEANDRO, CA 94577 ATTN CHARLES H. STALLINGS ATTN JOHN H. HUNTINGTON

POWER PHYSICS CORPORATION 542 INDUSTRIAL WAY WEST PO BOX 626 EATONTOWN, NJ 07724 ATTN MITCHELL BAKER

R & D ASSOCIATES PO BOX 9695 MARINA DEL REY, CA 90291 ATTN S. CLAY ROGERS

RAYTHEON COMPANY
HARTWELL ROAD
BEDFORD, MA 01730
ATTN GAJANAN H. JOSHI, RADAR SYS LAB

RATHEON COMPANY 528 BOSTON POST ROAD SUDBURY, MA 01776 ATTN HAROLD L. FLESCHER

RCA CORPORATION
GOVERNMENT & COMMERCIAL SYSTEMS
ASTRO ELECTRONICS DIVISION
PO BOX 800, LOCUST CORNER
PRINCETON, NJ 08540
ATTN GEORGE J. BRUCKER

RCA CORPORATION
DAVID SARNOFF RESEARCH CENTER
W. WINDSOR TWP
201 WASHINGTON ROAD, PO BOX 432
PRINCETON, NJ 08540
ATTN K. H. ZAININGER

RCA CORPORATION CAMBEN COMPLEX FRONT & COOPER STREETS CAMBEN, NJ 08012 ATTN E. VAN KEUREN, 13-5-2

RENSSELAER POLYTECHNIC INSTITUTE PO BOX 965 TROY, NY 12181 ATTN RONALD J. GUTMANN

RESEARCH TRIANGLE INSTITUTE
PO BOX 12194
RESEARCH TRIANGLE PARK, NC 27709
ATTN SEC OFFICER FOR
ENG DIV, MAYRANT SIMONS, JR.

ROCKWELL INTERNATIONAL CORPORATION 3370 MIROLOMA AVENUE ANAHEIM, CA 92803 ATTN K. F. HULL ATTN JAMES E. BELL, HA10 ATTN DONALD J. STEVENS, FA70 ATTN GEORGE C. MESSENGER, FB61 ATTN N. J. RUDIE. FA53

ROCKWELL INTERNATIONAL CORPORATION 5701 WEST IMPERIAL HIGHWAY LOS ANGELES, CA 90009 ATTN T. B. YATES

ROCKWELL INTERNATIONAL CORPORATION ELECTRONICS OPERATIONS COLLINS RADIO GROUP 5225 C. AVENUE NE CEDAR RAPIDS, IA 52406 ATTN ALAN A. LANGENFELD ATTN DENNIS SUTHERLAND ATTN MILDRED A. BLAIR

SANDERS ASSOCIATES, INC. 95 CANAL STREET NASHUA, NH 03060 ATTN M. L. AITEL, NCA 1-3236

SCIENCE APPLICATIONS, INC. PO BOX 2351 LA JOLLA, CA 92038 ATTN LARRY SCOTT ATTN J. ROBERT BEYSTER

SCIENCE APPLICATIONS, INC. HUNTSVILLE DIVISION 2109 W. CLINTON AVENUE SUITE 700 HUNTSVILLE, AL 35805 ATTN NOEL R. BYRN SCIENCE APPLICATIONS, INC. 2680 HANOVER STREET PALO ALTO, CA 94303 ATTN CHARLES STEVENS

SCIENCE APPLICATIONS, INC 8400 WESTPARK DRIVE MCLEAN, VA 22101 ATTN WILLIAM L. CHADSEY

SIMULATION PHYSICS, INC. 41 "B" STREET BURLINGTON, MA 01803 ATTN ROGER G. LITTLE

SINGER COMPANY, THE 1150 MC BRIDE AVENUE LITTLE FALLS, NJ 07424 ATTN IRWIN GOLDMAN, ENG MANAGEMENT

SINGER COMPANY (DATA SYSTEMS), THE 150 TOTOWA ROAD WAYNE, NJ 07470 ATTN TECH INFO CENTER

SPERRY FLIGHT SYSTEMS DIVISION SPERRY RAND CORPORATION P.O. BOX 21111 PHOENIX, AZ 85036 ATTN D. ANDREW SCHOW

SPERRY RAND CORPORATION UNIVAC DIVISION DEFENSE SYSTEMS DIVISION P.O. BOX 3525 MAIL STATION 1931 ST. PAUL, MN 55101 ATTN JAMES A. INDA, MS 41T25

SPERRY RAND CORPORATION
SPERRY DIVISION
SPERRY GYROSCOPE DIVISION
SPERRY SYSTEMS MANAGEMENT DIVISION
MARCUS AVENUE
GREAT NECK, NY 11020
ATTN PAUL MARRAFFINO
ATTN CHARLES L, CRAIG EV

STANFORD RESEARCH INSTITUTE 333 RAVENSWOOD AVENUE MENLO PARK, CA 94025 ATTN MR. PHILIP DOLAN

STANFORD RESEARCH INSTITUTE 306 WYNN DRIVE, N. W. HUNTSVILLE, AL 35805 ATTN MACPHERSON MORGAN

SUNDSTRAND CORPORATION
4751 HARRISON AVENUE
ROCKFORD, IL 61101
ATTN DEPT 763SW, CURTIS WHITE

SYSTRON-DONNER CORPORATION 1090 SAN MIGUEL ROAD CONCORD, CA 94518 ATTN HAROLD D. MORRIS ATTN GORDON B. DEAN

TEXAS INSTRUMENTS, INC. P.O. BOX 5474 DALLAS, TX 75222 ATTN DONALD J. MANUS, MS 72 TEXAS TECH UNIVERSITY
PO BOX 5404 NORTH COLLEGE STATION
LUBBOCK, TX 79417
ATTN TRAVIS L. SIMPSON

TRW SYSTEMS GROUP
ONE SPACE PARK
REDONDO BEACH, CA 90278
ATTN TECH INFO CENTER/S-1930
ATTN O. E. ADAMS, R1-1144
ATTN R. K. PLEBUCH, R1-2078
ATTN ROBERT M. WEBB, R1-2410
ATTN H. H. HOLLOWAY, R1-2036

TRW SYSTEMS GROUP SAN BERNARDINO OPERATIONS PO BOX 1310 SAN BERNARDINO, CA 92402 ATTN EARL W. ALLEN ATTN F. B. FAY, 527/710 ATTN R. KITTER

UNITED TECHNOLOGIES CORPORATION HAMILTON STANDARD DIVISION BRADLEY INTERNATIONAL AIRPORT WINDSOR LOCKS, CT 06069 ATTN RAYMOND G. GIGUERE

WESTINGHOUSE ELECTRIC CORPORATION
DEFENSE AND ELECTRONIC SYSTEMS CENTER
P.O. BOX 1693
FRIENDSHIF INTERNATIONAL AIRPORT
BALTIMORE, MD 21203
ATTN HENRY P. KALAPACA, MS 3525

HARRY DIAMOND LABORATORIES
ATTN MCGREGOR, THOMAS, COL, COMMANDING
OFFICER/FLYER, I.N./LANDIS, P.E./
SOMMER, H./OSWALD, R. B.
ATTN CARTER, W.W., DR., ACTING TECHNICAL
DIRECTOR/MARCUS, S.M.

ATTN CARTER, W.W., DR., ACT DIRECTOR/MARCUS, S.M.
ATTN KIMMEL, S., IO
ATTN CHIEF, 0021
ATTN CHIEF, 0022
ATTN CHIEF, LAB 100
ATTN CHIEF, LAB 200
ATTN CHIEF, LAB 400
ATTN CHIEF, LAB 400
ATTN CHIEF, LAB 500
ATTN CHIEF, LAB 600
ATTN CHIEF, LAB 600
ATTN CHIEF, DIV 700
ATTN CHIEF, DIV 700
ATTN CHIEF, LAB 900
ATTN CHIEF, LAB 900
ATTN CHIEF, LAB 1000

ATTN CHAIRMAN, EDITORIAL COMMITTEE ATTN CHIEF, 047 ATTN TECH REPORTS, 013 ATTN FATENT LAW BRANCH, 071 ATTN MCLAUGHLIN, P.W., 741

ATTN LANHAM, C., PROGRAM & PLANS OFFICE ATTN TOMPKINS, J. E., 230 ATTN MILETTA, J. R., 240 ATTN ROSADO, JOHN A., 240 ATTN CALDWELL, PAUL A., 290 ATTN WIMENITZ, F. N., 0024 ATTN SOKOLOSKI, M. S., 0024

ATTN SOKOLOSKI, M. S., 0024 ATTN MCGARRITY, J. M., 280 ATTN HALPIN, J. J., 280 ATTN BOESCH, H. E., JR. (15 COPIES)

ATTN MCLEAN, F. B., 280 ATTN WINOKUR, P. S., 280

The state of the s